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Applicant: KIMBERLY-CLARK CORPORATION 401 North Lake Street Neenah, Wisconsin 54956-0349(US)

Inventor: Sasse, Philip Anthony 5020 Morton Ferry Circle Alpharetta, Georgia 30202(US) Inventor: Durrance, Debra Hartley 581 Sunfield Drive Lilburn, Georgia 30247(US) Inventor: Taylor, Glenn Alfred Eichenweg 19 CH-4565 Obergerlaafingen(US)

PARTNER
Patentanwälte Flüggenstrasse 13
W-8000 München 19 (DE)

- <sup>54</sup> Polymeric compositions and use of same.
- The properties of (meth)acrylic ester/(meth)acrylic acid copolymer webs, such as films and nonwovens, are improved by blending the ester/acid copolymer with a modified polyolefin, such as an ethylene/(meth)acrylic acid copolymer and/or a styrene/butadiene block co polymer, and a polyether block amide copolymer. The copolymer blend provides a polymeric material which is useful for making personal care products such as diapers and feminine pads in that it can be made water-soluble in alkaline solutions while exhibiting other properties which are necessary for adequate product performance. Such materials are also useful for packaging products, such as film for plastic bags and the like.

In the production of personal care products, a number of different components and materials are required to construct the products. In the case of diaper manufacture, for example, thes components include a backing material, which is typically a film, and an inner liner, which is typically a nonwoven web. Also, composite structures of synthetic and natural fibers have utility as absorbent media in a variety of personal care products. These various synthetic components are typically made from thermoplastic polymers such as polyethylene or polypropylene. However, with a greater emphasis being placed on protecting the environment today, there is a need to develop materials which are more compatible with existing and developing waste disposal technologies while still delivering the performance consumers have come to expect.

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Copolymers of (meth)acrylate esters and (meth)acrylic acid are of environmental interest because of their solubility in alkaline solutions and their compostability due to eventual disintegration upon prolonged exposure to moisture, even though they are relatively hydrophobic. Unfortunately, the physical properties which make these materials desirable from an environmental standpoint can make them unsuitable for personal care products. In particular, films made from these polymers in contact with synthetic urine for a period greater than one hour will become hydrated, weak and sticky. This is obviously unacceptable for use in diapers, for example. Films made from these polymers also suffer from a lack of toughness and tear resistance. Films and nonwovens made from these copolymers are somewhat sticky and tend to stick or "block" in roll form. In addition, dimensional stability and aesthetic properties of these materials are also poor.

Therefore there is a need for copolymers of (meth)acrylate esters and (meth)acrylic acid which have modified properties suitable for use as components in personal care products such as diapers, feminine pads, incontinence garments and the like, and packaging products such as film for plastic bags and the like.

This object is solved by the composition of matter of independent claim 1, the polymeric web 11 of independent claim 11, the film of independent claim 13, and the absorbent articles of independent claim 26. Further advantageous features, aspects and details of the invention are evident from the dependent claims, the description, and the examples. The claims are intended to be understood as a first non-limiting approach of defining the invention in general terms.

The invention also provides polymeric web compositions having improved alkaline solubility.

The invention is based on the finding that copolymers of (meth)acrylate esters and/or (meth)-

acrylic acid can be modified with additional materials to provide polymeric webs having improved properties suitable for use in personal care products. More specifically, it has been discovered that modified polyolefins, such as copolymers of ethylene and (meth)acrylic acid and/or block copolymers of styrene and butadiene, and polyether block amide copolymers, when blended with these ester/acid copolymers in controlled ratios, impart increased resistance to moisture for webs made therefrom while retaining the desired selectve alkaline solubility. In addition, the tackiness of such webs is decreased, and dimensional stability and hand are markedly improved.

Hence, in one aspect, the invention resides in a composition of matter comprising a polymeric blend of three major components, namely: from about 50 to about 90 weight percent, preferably from about 60 to about 90 weight percent, and most preferably from about 70 to about 90 weight percent, of a (meth)acrylate ester/(meth)acrylic acid copolymer; from about 5 to about 30 weight percent, preferably from about 10 to about 15 weight percent, of a modified polyolefin such as a copolymer of ethylene and (meth)acrylic acid and/or a block copolymer of styrene and butadiene; and from about 1 to about 10 weight percent of a polyether block amide copolymer. Because the alkaline solubility of these compositions decreases with increasing amounts of the polyether block amide copolymer and becomes very low at polyether block amide copolymer levels above about 5 weight percent, a preferred composition contains from about 1 to about 5 weight percent, most preferably from about 3 to about 5 weight percent, of the polyether block amide copolymer. However, compositions with greater amounts are still useful for applications requiring disintegration upon prolonged exposure to moisture (e.g. in composting) rather than rapid dissolution in alkaline media.

In another aspect, the invention resides in a polymeric web comprising a blend of from about 50 to about 90 weight percent, preferably from about 60 to about 90 weight percent, and most preferably from about 70 to about 90 weight percent, of a (meth)acrylate ester/(meth)acrylic acid copolymer, from about 5 to about 30 weight percent, preferably from about 10 to about 15 weight percent, of a modified polyolefin such as a copolymer of ethylene and (meth)acrylic acid and/or a block copolymer of styrene and butadiene, and from about 1 to about 10 weight percent, preferably from about 1 to about 5 weight percent, and most preferably from about 3 to about 5 weight percent, of a polyether block amide copolymer. The relative proportions of the three major components will depend upon the desired properties of

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the product into which they ar to b made. For us in making films, for example, it is preferred that the copolymer blend contain from about 50 to about 90 weight perc nt of an acrylate/methacrylic acid copolymer or blend of ethyl acrylate/(meth)acrylic acid copolymers, from about 5 to about 15 weight percent of an ethylene/acrylic acid copolymer, and from about 3 to about 8 weight percent of a polyether block amide copolymer. For use in making nonwoven webs, such as spunbonded webs, it is preferred that the copolymer blend contain from about 70 to about 80 weight percent of an acrylate/methacrylic acid copolymer, from about 8 to about 15 weight percent of a modified polyolefin such as an ethylene/(meth)acrylic acid copolymer and/or a styrene/butadiene block copolymer, and from about 3 to about 5 weight percent of a polyether block amide copolymer.

In a further aspect, the invention resides in an absorbent article having an outer cover, an absorbent core, and an inner liner, wherein any or all of said outer cover, absorbent core and inner liner comprise the webs (nonwoven or film) described herein. Such absorbent articles particularly include diapers and sanitary napkins.

The (meth)acrylate ester/(meth)acrylic acid copolymer serves as the base material for these blends. This base material is preferably solely a pure copolymer, but can be modified with other monomers such as styrene to improve heat and water resistance. Hence for purposes herein, "-(meth)acrylate ester/(meth)acrylic acid copolymer" sometimes includes modified (meth)acrylate ester/-(meth)acrylic acid copolymers and blends of the pure copolymer with such modified copolymers. One preferred composition comprises ethyl acrylate and methacrylic acid as the major constituents with weight ratios of these two comonomers ranging from about 2:1 to about 4:1 (ester to acid). Another suitable composition comprises copolymers of ethyl acrylate and acrylic acid in the same weight ratio, with a comonomer weight ratio of about 3:1 (ester to acid) being preferred. Still another suitable composition comprises copolymers of methyl acrylate and methacrylic acid, preferably having a comonomer weight ratio of about 4:1 (ester to acid). The acrylate/methacrylic acid copolymer that is most preferred has a weight ratio of the comonomers from about 3.5:1 to about 4:1 (ester to acid). The weight average molecular weight of the most preferred ethyl acrylate/methacrylic acid copolymer is about 150,000, with a melt flow rate of about 7 grams per 10 minutes, as measured at 170°C using a 2160 g weight and a 2.1 mm by 8 mm capillary. Clearly, however, many similar copolymers can be prepared that will provide similar attributes and can be substitut d for the most pr ferred copolymer in thes compositions. For example, any other (meth)acrylate ester derived from an alcohol having from 1 to 18 carbon atoms can be substituted for all or part of the ethyl acrylate. Such substitutions can lead to enhancement of particular properties for specific material applications. Although not limited to one method of manufacture, a method of making such copolymers is described in U.S. Patent No. 4,870,148 to RB Kunststoffpatent-Verwertugs AG and Belland AG, both of Switzerland, issued September 26, 1989, which is herein incorporated by reference. Such copolymers are commercially available from Belland AG, and the most preferred copolymer is available as product codes GBC 2620 (ethyl acrylate/methacrylic acid copolymer, GBC 2518 (30 weight percent styrene-modified terpolymer), and GBC 1620 (methyl acrylate/methacrylic acid copolymer).

The ethylene/(meth)acrylic acid copolymers to be used as the modified polyolefins of these high melt index dispersible blends are ethylene/acrylic acid copolymers that are typically 20 percent acrylic acid by weight and 300 to 1300 in melt index. They are dispersible in alkaline water without emulsifiers, but are water-resistant in the acid form. Other suitable ethylene/acrylic acid copolymers which are available contain 3 to 9.5 percent acrylic acid by weight and have a melt index from 2.5 to 13.5. Such copolymers are commercially available from Dow Chemical Company, Midland, Michigan under the tradename PRIMACOR@, with the most preferred copolymers being those with product code numbers ranging from "5980" to "5990". They are produced by the free radical, high pressure copolymerization of ethylene and acrylic acid in a process similar to that used for producing low-density polyethylene. Also suitable for use in these blends are ethylene/ methacrylic acid copolymers. Such copolymers are commercially available from E.I. du Pont, Wilmington. Delaware, under the tradename NUCREL@, with grades ranging from 4 to 12 percent methacrylic acid by weight and from 2.5 to 13.5 in melt index.

The styrene/butadiene block copolymers to be used as the modified polyolefin component of these blends are synthetic rubbers typically containing from about 20 to about 40 weight percent styrene and having a melt index of from about 1 to about 20. Specific commercially available materials include CARIFLEX@ TR-1102 or KRATON D-1102 (Shell Chemical Company) and K-Resin KR-03 (British Petroleum Company). Both of these materials provide th same flex properties as the PRIMACOR material described above, but are less soluble and form a hazy blend rather than a clear

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The polyether block amid copolymers that are useful in the blends of this invention are thermoplastic elastomers consisting of flexible polyether segments connected with rigid polyamide segments into linear block copolymers. The most suitable polyether block amide copolymers are commercially available from Atochem under the tradename PEBAX@ with grade designations 2533, 3533, 4533 and 5533. Of these grades, the most preferred are the grades designated 2533 and 3533.

In all aspects of this invention, combinations or blends of more than one species within one or more of the three major components can be used to prepare the compositions useful for purposes of this invention and are considered to be equivalents for all purposes. Hence for purposes of convenience, unless otherwise specified, any reference to a copolymer is to be interpreted to include one or more copolymers within the particular copolymer genus.

Blends of the three major components can be prepared by mixing the desired weight ratio of the polymer pellets and blending them using any standard equipment commonly used for blending thermoplastic polymers under conditions of heat and high shear. These include the Banbury@ type of intensive production mixer (Farrel Corp, Ansonia, CT) and both single-and twin-screw compounding extruders, which can utilize high-shear mixing screws, co-rotating kneading blocks, etc.

In addition to blends containing the three above-mentioned major components, other components can be added to further enhance the properties of the resulting material. For example, polyethylene glycol can be added to lower the melt viscosity of these copolymers to a range suitable for meltblown or meltsprayed nonwovens and also improve the flexibility of the resulting webs. The amount of polyethylene glycol added to the copolymer blend can be from about 5 to about 20 weight percent, based on the total weight of the final blend, and a preferred range is from about 10 to about 15 weight percent. Suitable polyethylene glycols are available commercially from Union Carbide Corporation, Danbury, Connecticut, under the tradename CARBOWAX@; most suitable are product code numbers "3350" and "8000."

Polyethylene can also be added to blends containing the above-mentioned copolymers to improve the softness of the resulting nonwoven webs. The amount of polyethylene added to the copolymer blend can be from about 5 to about 15 weight percent, based on the total weight of the blend. The polyethylene grade must be selected so that the final blend has a melt index suitable for the nonwoven process to be used. Suitable fiber grade

poly thylenes ar available commercially from Dow Chemical Company, Midland, Michigan, under th tradename ASPUN@. Product code number "6811" is most suitable for blends for spunbond nonwovens and product code numbers "6806" and "6814" are most suitable for blends for meltblown or meltsprayed nonwovens.

Still further improvements to the properties of the webs of this invention, particularly films, can be made by adding certain fillers such as fumed silica, calcium carbonate or talc. Various particulate fillers have been shown to reduce blocking, noise and gloss in the films. Such materials can be added in amounts of from about 2 percent to about 20 weight percent, based on the total weight of the blend. Processing characteristics of the blends for both films and nonwovens can be improved by the incorporation of lubricants or slip agents into the blends. Additives of other types normally used in polymer blends can also be incorporated to provide specific properties as needed, such as antistatic agents, pigments or other colorants, and the like. All of these additive types are generally used in small amounts, usually less than 5 percent.

Films of the copolymer blends can be prepared by extrusion of the blend through a linear film die, allowing the film to attenuate under its own weight into a nip between two rolls (cast film). Alternatively, an annular die can be used to produce a polymeric tube, which can be attenuated by an air stream to form a film "bubble" (blown film).

Nonwoven webs of the copolymer blends can be prepared by extrusion of the blend through a plurality of capillaries, producing a series of filaments. These filaments can be quenched and then attenuated into fibers by an accelerating gas stream. The fibers can be collected on a moving surface, where they are deposited by the gas stream in a random fashion. Passing the resulting batt through a pair of heated rolls bonds the fibers together into an integral web. Alternatively, a hot gas stream may be used to attenuate and break the filaments in the molten state. These discontinuous fibers can be collected on a moving surface, where they will lay down in a random, entangled manner, producing an integral web. Suitable nonwoven webs include, without limitation, meltblown webs, spunbonded webs, and coform webs (meltblown webs in which a second fiber source, such as cellulose fibers, is blown into the primary meltblown fiber stream prior to deposition onto the collecting surface). All of such webs are known in the nonwovens art.

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#### **EXAMPLES**

# Example 1: Preparation of Copolymer Blend

A blend in accordance with this invention was prepared containing 80 percent by weight of Belland's copolymer material designated GBC 2620 AC. This material is reported to be a copolymer of ethyl acrylate and methacrylic acid in a ratio of 4:1 by weight, respectively, containing 1 percent by weight of an antioxidant (Ciba Geigy, Irganox 1076, octadecyl 3-(3,5-di-t-butyl-4-hydroxyphenyl) propionate) and 3 percent by weight of Hostalube@ FA-1 (Hoescht Celanese, mixed amide of ethylenediamine with palmitic and stearic acids). This material was compounded with 11 percent by weight of Primacor@ 5981 (Dow, ethylene-acrylic acid copolymer), 5 percent by weight of PEBAX 2533SA, 2 percent by weight of titanium dioxide pigment, and 2 percent by weight of erucamide as a slip agent.

The ingredients were first mixed thoroughly by means of a drum tumbler, and then transferred to an AccuRate@ volumetric feeder. The mixture was metered into a Werner & Pfleiderer 30-mm twinscrew compounding extruder for blending. Extruder zones 1 through 6 were set with the following temperature profile (in \*C): 100, 101, 110, 121, 140, 145; with the lowest temperature being at the feed zone adjacent to the throat of the extruder where the mixture enters and the highest temperature being at the die from which the molten blend exits.

The extruder screw configuration used was a vented, two-stage mixing screw combination of a type typically used for preparing polyolefin blends. Extruder throughput was maintained at ca. 22.7 kg/hour (ca. 50 pounds/hour). The molten blend was taken off from a four-strand die into a water bath with a dip length of six feet, then through two air knives to remove as much water as possible. The pellets were ejected from a rotating-knife pelletizer directly into a spin dryer to remove the remaining surface moisture before screening and packaging.

### Example 2: Preparation of Copolymer Blend Film

A copolymer blend containing 73 weight percent GBC 2620 WB (the WB grade contains titanium dioxide pigment), 10 weight percent Primacor@ 5990, 5 weight percent PEBAX, 2 weight percent erucamide, and 10% talc was blended as in Example 1. The polymer blend was then processed at a rate of 115 pounds/hour through a single-screw extruder with a 8.9 cm (3.5 inch) diameter, internally-cooled screw (3:1 compression ratio, 24:1 L/D). A flat t mperature profile

of 127°C (260°F) was us d in all extruder zones, transfer piping and in the film die. Film was produced using a six-inch diameter annular die with a die gap of 0.1 cm (0.042 inches). A bubble was generated with a diameter of 35.5 - 45.7 cm (14-18 inches), producing film that varied from .0028 to .0036 cm (0.0011 to 0.0014 inches) in gauge.

Ultimate tensile strengths were determined to be 1356 grams/inch at 255% elongation (machine direction) and 1000g/2.54 cm (1000 grams/inch) at 285% elongation (cross direction). A similarly-prepared film of 100% GBC 2620 WB with the same gauge of .0028 cm (0.0011 inch) had ultimate tensile strengths of 2647 g/2.54 cm (2647 grams/inch) at 224% elongation (machine direction) and 1274 g/2.54 cm (1274 grams/inch) at 365% elongation (cross direction). The films prepared from the described blend show less isotropy, less tendency to block, better moisture resistance, and better tear resistance than the film prepared from GBC 2620 WB alone.

# Example 3: Preparation of Copolymer Blend Non-woven Web

A copolymer blend of this invention was prepared as described in Example 1. A nonwoven web was prepared from this blend by means of a spunbond process. The polymer blend pellets were introduced into a single-screw extruder having a screw configuration of a type normally used for extrusion of polyolefins. The molten polymer, at a melt temperature of 171° C (340° F), was conveyed from the extruder through a metering pump to a fiber-spinning die. Molten polymer strands exiting the die were quenched, attenuated and formed into a web by controlled streams of high-velocity air. The resulting web was carried by a forming wire through compaction and bonding sections to form a point-bonded spunbond nonwoven.

Webs formed in this manner are dimensionally stable (less than 5% shrinkage) and possess acceptable drape, flexibility and softness. Webs prepared by the same process from GBC 2620 with no additives suffered severe shrinkage (up to 50%), distortion, and increasing rigidity over several weeks. Moisture was found to accelerate these changes.

#### Claims

 A composition of matter comprising a blend of from 50 to 90 weight percent of a copolymer of a (meth)acrylate ester and (meth)acrylic acid, from 5 to 30 weight percent of a copolymer of ethylene and (meth)acrylic acid, or of a styrene/butadiene block copolymers and from 1 to 10 weight percent of a polyether block

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amid copolymer.

- The composition of Claim 1 wherein the amount of th polyether block amide copolymer is from 3 to 8 weight percent, preferably from 3 to 5 weight percent.
- The composition of Claim 1 or 2 further comprising from 5 to 15 weight percent polyethylene.
- The composition of any one of the preceding Claims wherein the (meth)acrylate ester is ethyl acrylate.
- The composition of Claim 4 wherein the ethyl acrylate is copolymerized with methacrylic acid.
- 6. The composition of any one of the preceding Claims wherein the (meth)acrylic acid of the ethylene/(meth)acrylic acid copolymer is acrylic acid.
- 7. The composition of Claim 6 wherein the copolymer of ethyl acrylate and methacrylic acid comprises about 80 weight percent ethyl acrylate moieties and about 20 weight percent methacrylic acid moieties.
- 8. The composition of Claim 6 wherein the copolymer of ethylene and acrylic acid comprises about 80 weight percent ethylene moieties and about 20 weight percent acrylic acid moieties.
- 9. The composition of Claim 6 comprising from 70 to 90 weight percent of a copolymer of ethyl acrylate and methacrylic acid, from 10 to 15 weight percent of a copolymer of ethylene and (meth)acrylic acid,and from 3 to 5 weight percent of a polyether block amide copolymer.
- 10. The composition of Claim 6 comprising from 60 to 90 weight percent of a copolymer of ethyl acrylate and methacrylic acid, from 5 to 10 weight percent of a copolymer of ethylene and (meth)acrylic acid, and from 3 to 5 weight percent of a polyether block amide copolymer.
- 11. A polymeric web comprising a blend as defined in any one of the preceding claims whereby the web is proferable nonwoven.
- 12. The web of Claim 11 wherein the amount of the polyether block amide copolymer is from I to 5 weight percent.

- A film comprising a blend according to any one of claims 1 to 10.
- 14. The film of claims 13 comprising of from 60 to 90 weight percent of a copolymer of ethyl acrylate and methacrylic acid, from 5 to 10 weight percent of a copolymer of ethylene and (meth)acrylic acid, and from 3 to 5 weight percent of a polyether block amide copolymer, wherein said copolymer of ethyl acrylate and methacrylic acid comprises about 80 weight percent ethyl acrylate moieties and about 20 weight percent methacrylic acid moieties and wherein said copolymer of ethylene and (meth)acrylic acid contains about 80 weight percent ethylene moieties and about 20 weight percent acrylic acid moieties.
- 15. The film of Claim 14 further comprising from 5 to 15 weight percent polyethylene.
- 16. The nonwoven web especially of claim 11 comprising a blend of from 70 to 90 weight percent of a copolymer of ethyl acrylate and methacrylic acid, from 10 to 15 weight percent of a copolymer of ethylene and (meth)acrylic acid, and from 3 to 5 weight percent of a polyether block amide copolymer, wherein said copolymer of ethyl acrylate and methacrylic acid comprises about 80 weight percent ethyl acrylate moieties and about 10 weight percent methacrylic acid moieties and wherein said copolymer of ethylene and acrylic acid comprises about 80 weight percent ethylene moieties and about 20 weight percent acrylic acid moieties.
- The nonwoven web of Claim 16 further comprising from 5 to 15 weight percent polyethylene.
- 18. The nonwoven web of Claim 16 or 17 comprising a blend of from 70 to about 80 weight percent of a copolymer of ethyl acrylate and methacrylic acid and from 10 to 15 weight percent of a copolymer of ethylene and acrylic acid.
- 19. The nonwoven web of any one of claims 16 to 18 further comprising from 5 to 20 weight percent of polyethylene glycol.
- 20. The nonwoven web of any one of Claims 16 to 18 further comprising from 10 to 15 weight percent of polyethylene glycol.
- 21. The nonwoven web of Claim 16 comprising a blend of about 80 weight percent of the

copolymer of thyl acrylate and methacrylic acid, about 10 weight percent of th copolym r of ethylene and acrylic acid, and about 10 weight percent poly thylene glycol.

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- 22. The nonwoven web of Claim 16 further comprising from about 5 to about 15 weight percent polycaprolactone.
- **23.** The nonwoven web of any one of Claims 16 to 22 wherein said web is a meltblown web.
- 24. The nonwoven web of any one of Claims 16 to 22 wherein said web is a spunbonded web.

25. The nonwoven web of any one of Claims 16 to 22 wherein said web is a coform web.

- 26. An absorbent article comprising an outer cover, an absorbent core and an inner liner, wherein either of said outer cover or said inner liner is a nonwoven web comprising a blend as defined in any one of claims 1 to 10.
- 27. The article of Claim 26 wherein the amount of the polyether block amide copolymer is from 1 to 5 weight percent.
- 28. The article of Claim 26 or 27 comprising from 70 to 90 weight percent of a copolymer of ethyl acrylate and methacrylic acid and from 10 to 15 weight percent of a copolymer of ethylene and acrylic acid.
- 29. The article of Claim 26 or 27 comprising from 80 to 90 weight percent of a copolymer of ethyl acrylate and methacrylic acid and from 10 to 20 weight percent of a copolymer of ethylene and acrylic acid.
- 30. The absorbent article of any one of claims 26 to 29, wherein said absorbent article is a diaper.
- 31. The absorbent article of any one of Claims 26 to 29, wherein said absorbent article is a sanitary napkin.

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Application Number

EP 92 11 3489

ategory	Citation of document with indica of relevant passag	ation, where appropriate,	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)	
	EP-A-0 352 624 (MITSUI * page 4, line 11 - 1 * page 7, line 46 - page 7 * claims 1-2 *	BISHI RAYON CO)	1-2	C08L33/06 C08J5/18 D04H1/56 A61L15/00 //(C08L33/06, 33:02,53:00) (C08L33/06, 23:08,53:00)	
				TECHNICAL FIELDS SEARCHED (Int. Cl.5) CO8L CO8J DO4H A61L	
	The present search report has bee	n drawn up for all claims	_		
	Place of search	Date of complation of the search 03 DECEMBER 1992		Examiner SIEMENS T.	
X : p	THE HAGUE  CATEGORY OF CITED DOCUMENT articularly relevant if taken alone articularly relevant if conshined with anoth	T: theory or print E: earlier patent after the filli D: document of	or principle underlying the invention patent document, but published on, or e filing date of cited in the application		
A: t O: c	ocument of the same category echnological background ion-written disclosure atermediate document	L : document cit	L: document cited for other reasons  à: member of the same patent family, corresponding		